

<u>NEWS 1</u>			Web Page for STN Seminar Schedule - N. America
<u>NEWS 2</u>	MAR 31		IFICDB, IFIPAT, and IFIUDB enhanced with new custom IPC display formats
<u>NEWS 3</u>	MAR 31		CAS REGISTRY enhanced with additional experimental spectra
<u>NEWS 4</u>	MAR 31		CA/CAplus and CASREACT patent number format for U.S. applications updated
<u>NEWS 5</u>	MAR 31		LPCI now available as a replacement to LDPCI
<u>NEWS 6</u>	MAR 31		EMBASE, EMBAL, and LEMBASE reloaded with enhancements
<u>NEWS 7</u>	APR 04		STN AnaVist, Version 1, to be discontinued
<u>NEWS 8</u>	APR 15		WPIDS, WPINDEX, and WPIX enhanced with new predefined hit display formats
<u>NEWS 9</u>	APR 28		EMBASE Controlled Term thesaurus enhanced
<u>NEWS 10</u>	APR 28		IMSRESEARCH reloaded with enhancements
<u>NEWS 11</u>	MAY 30		INPAFAMDB now available on STN for patent family searching
<u>NEWS 12</u>	MAY 30		DGENE, PCTGEN, and USGENE enhanced with new homology sequence search option
<u>NEWS 13</u>	JUN 06		EPFULL enhanced with 260,000 English abstracts
<u>NEWS 14</u>	JUN 06		KOREAPAT updated with 41,000 documents
<u>NEWS 15</u>	JUN 13		USPATFULL and USPAT2 updated with 11-character patent numbers for U.S. applications
<u>NEWS 16</u>	JUN 19		CAS REGISTRY includes selected substances from web-based collections
<u>NEWS 17</u>	JUN 25		CA/CAplus and USPAT databases updated with IPC reclassification data
<u>NEWS 18</u>	JUN 30		AEROSPACE enhanced with more than 1 million U.S. patent records
<u>NEWS 19</u>	JUN 30		EMBASE, EMBAL, and LEMBASE updated with additional options to display authors and affiliated organizations
<u>NEWS 20</u>	JUN 30		STN on the Web enhanced with new STN AnaVist Assistant and BLAST plug-in
<u>NEWS 21</u>	JUN 30		STN AnaVist enhanced with database content from EPFULL
<u>NEWS 22</u>	JUL 28		CA/CAplus patent coverage enhanced
<u>NEWS 23</u>	JUL 28		EPFULL enhanced with additional legal status information from the epoline Register
<u>NEWS 24</u>	JUL 28		IFICDB, IFIPAT, and IFIUDB reloaded with enhancements
<u>NEWS 25</u>	JUL 28		STN Viewer performance improved
<u>NEWS 26</u>	AUG 01		INPADOCDB and INPAFAMDB coverage enhanced

NEWS EXPRESS JUNE 27 08 CURRENT WINDOWS VERSION IS V8.3,
AND CURRENT DISCOVER FILE IS DATED 23 JUNE 2008.

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=> file caplus
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ENTRY SESSION

FULL ESTIMATED COST

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0.21

FILE 'CAPLUS' ENTERED AT 07:50:40 ON 04 AUG 2008
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FILE COVERS 1907 - 4 Aug 2008 VOL 149 ISS 6
 FILE LAST UPDATED: 3 Aug 2008 (20080803/ED)

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=> tetrafluorborate (s) (metal or li or lithium or na or sodium or k or potassium)
 TETRAFLUORBORATE IS NOT A RECOGNIZED COMMAND
 The previous command name entered was not recognized by the system.
 For a list of commands available to you in the current file, enter
"HELP COMMANDS" at an arrow prompt (=>).

=> s tetrafluorborate (s) (metal or li or lithium or na or sodium or k or potassium)
 28 TETRAFLUORBORATE
 1854028 METAL
 208489 LI
 345220 LITHIUM
 807162 NA
 1212655 SODIUM
 1470040 K
 688225 POTASSIUM
 L1 2 TETRAFLUORBORATE (S) (METAL OR LI OR LITHIUM OR NA OR SODIUM OR
 K OR POTASSIUM)

=> s tetrafluoroborate (s) (metal or li or lithium or na or sodium or k or potassium)
 25067 TETRAFLUOROBORATE
 1854028 METAL
 208489 LI
 345220 LITHIUM
 807162 NA
 1212655 SODIUM
 1470040 K
 688225 POTASSIUM
 L2 5225 TETRAFLUOROBORATE (S) (METAL OR LI OR LITHIUM OR NA OR SODIUM
 OR K OR POTASSIUM)

=> s bf4 (s) (metal or li or lithium or na or sodium or k or potassium)
 19010 BF4
 1854028 METAL
 208489 LI
 345220 LITHIUM
 807162 NA
 1212655 SODIUM

1470040 K
 688225 POTASSIUM
 L3 2137 BF4 (S) (METAL OR LI OR LITHIUM OR NA OR SODIUM OR K OR POTASSIU
 M)

=> s ?ammonium (s) (bf4 or tetrafluoroborate)
 550598 ?AMMONIUM
 19010 BF4
 7 TETRAFLUOROBORATE
 L4 7 ?AMMONIUM (A) (BF4 OR TETRAFLUOROBORATE)

=> s ?ammonium (a) (bf4 or tetrafluoroborate)
 550598 ?AMMONIUM
 19010 BF4
 25067 TETRAFLUOROBORATE
 L5 3706 ?AMMONIUM (A) (BF4 OR TETRAFLUOROBORATE)

=> s (l1 or l2) and (l3 ad l4)
 MISSING OPERATOR L3 AD
 The search profile that was entered contains terms or
 nested terms that are not separated by a logical operator.

=> s (l1 or l2) and (l3 or l4)
 L6 334 (L1 OR L2) AND (L3 OR L4)

=> s ?ammonium (s) (halide or fluoride or bromide or chloride or iodide)
 550598 ?AMMONIUM
 160802 HALIDE
 276080 FLUORIDE
 297824 BROMIDE
 1219594 CHLORIDE
 194356 IODIDE
 L7 128703 ?AMMONIUM (A) (HALIDE OR FLUORIDE OR BROMIDE OR CHLORIDE OR IODI
 DE)

=> s l5 and l7
 L8 413 L5 AND L7

=> s l6 and l7
 L9 8 L6 AND L7

=> d l-8 bib, ab

L9 ANSWER 1 OF 8 CAPLUS COPYRIGHT 2008 ACS on STN
 Full Text
 AN 2008:566491 CAPLUS
 DN 149:113149
 TI Surface tension of binary mixtures of imidazolium and ammonium based ionic
 liquids with alcohols, or water: Cation, anion effect
 AU Domanska, Urszula; Pobudkowska, Aneta; Rogalski, Marek
 CS Physical Chemistry Division, Faculty of Chemistry, Warsaw University of
 Technology, Warsaw, 00-664, Pol.
 SO Journal of Colloid and Interface Science (2008), 322(1), 342-350
 CODEN: JCISA5; ISSN: 0021-9797
 PB Elsevier
 DT Journal
 LA English
 AB The surface tensions were measured at atm. pressure, with use of a ring
 tensiometer, of a series of alc. solns. of closely related ionic liqs.:
 1-methyl-3-methylimidazolium methylsulfate, [MMIM][CH₃SO₄] in alc.
 (methanol, or ethanol, or 1-butanol at 298.15 K), 1-butyl-3-
 methylimidazolium methylsulfate, [BMIM][CH₃SO₄] in alc. (methanol, or
 ethanol, or 1-butanol at 298.15 K), 1-butyl-3-methylimidazolium
 octylsulfate, [BMIM][OcSO₄] in alc. (methanol, or 1-butanol at 298.15 K)
 and of 1-hexyloxyethyl-3-methylimidazolium tetrafluoroborate,

[C6H13OCH2MIM] [BF4], 1,3-dihexyloxymethylimidazolium tetrafluoroborate, [(C6H13OCH2)2IM] [BF4] in alc. (methanol, or 1-butanol, or 1-hexanol at 308.15 and 318.5 K) and hexyl(2-hydroxyethyl)dimethylammonium bromide, C6Br in 1-octanol at 298.15 K. The set of ammonium ionic liqs. of different cations and anions (C2Br, C2BF4, C2PF6, C2N(CN)2, C3Br, C4Br and C6Br) was chosen to show the influence of small amt. of the ammonium ionic liq. on the surface tension of water at 298.15 K. The influence of the cation, or anion alkyl chain length on the properties under study (densities and surface tension) was tested.

RE.CNT 43 THERE ARE 43 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L9 ANSWER 2 OF 8 CAPLUS COPYRIGHT 2008 ACS on STN

Full Text References

AN 2006:1005759 CAPLUS
DN 145:376914
TI Preparation of quaternary ammonium halides with low coloring and minimum content of amines and of quaternary salts therefrom
IN Nishimoto, Yoshihiro
PA Koei Chemical Co., Ltd., Japan
SO Jpn. Kokai Tokkyo Koho, 12pp.
CODEN: JKXXAF

DT Patent
LA Japanese

FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 2006257039	A	20060928	JP 2005-78439	20050318
PRAI JP 2005-78439		20050318		
OS CASREACT 145:376914; MARPAT 145:376914				
AB The halides Q+X- (Q = quaternary ammonium cation; X = halo), useful for intermediates of electrolytes or catalysts, are prep'd. by reacting tertiary amines with alkyl halides in the presence of basic alk. earth metal compds. [e.g., carbonates, (hydr)oxides, etc.]. The halides are ion exchanged with compds. M+A- [M+ = proton, metal, ammonium cation; A- = N(SO2CF3)-, BF4-, PF6-] to afford quaternary salts Q+A- (Q, A = the same as above). Thus, 30.0 g of 1-methylimidazole was reacted with 58.7 g 60.2% EtCl/MeCN in the presence of Ca(OH)2 at 90° to give 1-ethyl-3-methylimidazolium chloride (I) with APHA color 240 and amine content 150 ppm. Then, I was further reacted with NaBF4 in MeCN at 25° to give ion-exchanged product with APHA color 240 and amine content 100 ppm.				

L9 ANSWER 3 OF 8 CAPLUS COPYRIGHT 2008 ACS on STN

Full Text References

AN 2006:912889 CAPLUS
DN 145:293184
TI Improved process for preparation of alkoxyisilylmethyl isocyanurates by cyclocondensation of alkali metal isocyanates with chloromethylsilanes, catalyzed by tetraalkylammonium salts
IN Popp, Alfred; Stowischek, Klaus
PA Consortium fuer Elektrochemische Industrie G.m.b.H., Germany
SO Ger. Offen., 7pp.
CODEN: GWXXBX

DT Patent
LA German

FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
DE 102005009790	A1	20060907	DE 2005-102005009790	20050303
WO 2006092324	A1	20060908	WO 2006-EP1969	20060303
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE,				

GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ,
 LC, LK, LR, LS, LT, LU, LV, LY, MA, MD, MG, MK, MN, MW, MX, MZ,
 NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG,
 SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN,
 YU, ZA, ZM, ZW

RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE,
 IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ,
 CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH,
 GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY,
 KG, KZ, MD, RU, TJ, TM

EP 1853613 A1 20071114 EP 2006-707401 20060303

EP 1853613 B1 20080702

R: BE, DE, FR, GB, IT

CN 101133069 A 20080227 CN 2006-80006907 20070903

KR 2007108930 A 20071113 KR 2007-722539 20071002

PRAI DE 2005-102005009790 A 20050303

WO 2006-EP1969 W 20060303

OS CASREACT 145:293184; MARPAT 145:293184

AB Silylated 1,3,5-triazine-2,4,6-triones 1,3,5-[R1n(RO)3-nSiCH2]C3N3O3 [1, R = C4-15 organyl, COMe; R1 = H, (un)substituted C1-20 organyl, optionally having O, CO, CO2, OCO, OCO2, S, amino, imino, azo, phosphino groups in the main chain; n = 0-2; preferably R, R1 = Me, Et], useful as adhesives or additives for organosiloxanes and siloxane polymers (no data), were prepd. by cyclocondensation of chloromethylsilanes R1n(RO)3-nSiCH2Cl (same R, R1) with metal isocyanates M(OCN)m (M = alkali or alk. earth metal, m = 1, 2; preferably M = Na, K) in the presence of tetraalkylammonium salts R24N+X- (R2 = C1-20 organyl, optionally substituted by CN, OH, halo; X = Cl, Br, I, BF4-, BPh4; preferably R2 = Me, Et, Bu; R24N+ = PhCH2NBu3+; X = I-, BF4-) in dipolar org. solvent, preferably in DMF or DMF-contg. mixts., preferably having <200° b.p. at normal pressure, preferably in molar ratio M(OCN)m to R1n(RO)3-nSiCH2Cl of 1-1.2 at 0-200°, preferably at 80-100° in batch or continuous reactors. The use of tetraalkylammonium catalysts improves yields of 1 and reduces amts. of byproducts. In an example, 1,3,5-tris(methoxydimethylsilylmethyl)-1,3,5-triazine-2,4,6-trione (1, R = R1 = Me, n = 2) was prepd. by cyclocondensation of 0.33 mol of NaOCN with 0.3 mol of (MeO)Me2SiCH2Cl in 87.9 g of DMF in the presence of 1.6 mol% of Me4NI at 130° for 2 h with 87% yield; the yields of byproducts were less than 6.2%.

L9 ANSWER 4 OF 8 CAPLUS COPYRIGHT 2008 ACS on STN

Full Text

AN 2006:123472 CAPLUS

DN 144:182276

TI Cost-effective manufacture of (lower alkoxyalkyl)trimethylammonium salt ionic liquids

IN Amano, Yasutoshi; Noji, Kazuaki; Fujimoto, Masaki

PA Sanko Chemical Industry Co., Ltd., Japan

SO Jpn. Kokai Tokkyo Koho, 8 pp.

CODEN: JKXXAF

DT Patent

LA Japanese

FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
<u>PI JP 2006036652</u>	A	20060209	<u>JP 2004-214959</u>	20040722
<u>PRAI JP 2004-214959</u>		20040722		

OS MARPAT 144:182276

AB The ionic liqs., useful as electrolytes for elec. devices, are manufd. by treatment of NMe3 with lower alkoxyalkyl chlorides, followed by anion exchange of the resulting (lower alkoxyalkyl)trimethylammonium chlorides with YA [A = H+, alkali metal cation; Y = BF4-, CF3SO3-, CF3CO2-, (CnF2n+1MO2)2N-, PF6-, AsF6-, SbF6-, (CnF2n+1MO2)2N-; M = S, C; n = 1, 2]. Thus, NMe3 was treated with C3H8O(CH2)2Cl and anion-exchanged with NaBF4 to give C3H8O(CH2)2NMe3+BF4- with potential window from -3.0 V to 2.3 V.

L9 ANSWER 5 OF 8 CAPLUS COPYRIGHT 2008 ACS on STN

Full Text

AN 2003:537623 CAPLUS
 DN 139:390066
 TI An efficient synthesis for tetracyanoborates by sinter processes
 AU Bernhardt, E.; Finze, M.; Willner, H.
 CS Anorganische Chemie, Universitaet Duisburg, Germany
 SO Zeitschrift fuer Anorganische und Allgemeine Chemie (2003), 629(7-8), 1229-1234

CODEN: ZAACAB; ISSN: 0044-2313
 PB Wiley-VCH Verlag GmbH & Co. KGaA
 DT Journal
 LA German
 OS CASREACT 139:390066

AB A new efficient synthesis for tetracyanoborates in molar scale starting from the readily available reagents K[BF₄]⁻, LiCl, and KCN is presented. The tetracyanoborate, obtained after the sinter process and work up, is in contrast to the products of the reactions known so far a white solid without any impurities. The thermal behavior of the mixed cyanofluoroborates, K[BF_x(CN)_{4-x}] (x = 1-3), was investigated to solve the mechanism of the ligand exchange that takes place during the K[B(CN)₄] synthesis. Furthermore several metathesis reactions for the synthesis of different tetracyanoborates are described.

RE.CNT 24 THERE ARE 24 CITED REFERENCES AVAILABLE FOR THIS RECORD
 ALL CITATIONS AVAILABLE IN THE RE FORMAT

L9 ANSWER 6 OF 8 CAPLUS COPYRIGHT 2008 ACS on STN

Full Text

AN 2001:167660 CAPLUS
 DN 134:207547
 TI Preparation of stable (CF₃)₂N⁻ salts and their use in the synthesis of liquid crystals
 IN Heider, Udo; Schmidt, Michael; Sartori, Peter; Ignatyev, Nikolai; Kucheryna, Andrej
 PA Merck Patent G.m.b.H., Germany
 SO Eur. Pat. Appl., 11 pp.
 CODEN: EPXXDW

DT Patent
 LA German

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	<u>EP 1081129</u>	A2	20010307	<u>EP 2000-118126</u>	20000828
	<u>EP 1081129</u>	A3	20010321		
	<u>EP 1081129</u>	B1	20031022		
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO				
	<u>DE 19941566</u>	A1	20010308	<u>DE 1999-19941566</u>	19990901
	<u>TW 221832</u>	B	20041011	<u>TW 2000-89117401</u>	20000828
	<u>CA 2317284</u>	A1	20010301	<u>CA 2000-2317284</u>	20000830
	<u>BR 2000003885</u>	A	20010403	<u>BR 2000-3885</u>	20000830
	<u>CN 1286245</u>	A	20010307	<u>CN 2000-126092</u>	20000831
	<u>JP 2001122834</u>	A	20010508	<u>JP 2000-265452</u>	20000901
	<u>US 6582849</u>	B1	20030624	<u>US 2000-654519</u>	20000901
	<u>RU 2257376</u>	C2	20050727	<u>RU 2000-122754</u>	20000901
PRAI	<u>DE 1999-19941566</u>	A	19990901		

OS MARPAT 134:207547

AB Stable (CF₃)₂N⁻ salts [[[R₁(CR₂R₃)_k]_lAx]_y]K⁺-N(CF₃)₂ [A = N, P, P(:O), O, S, S(:O), SO₂, As, As(:O), Sb, Sb(:O); K = N, P, As, Sb, S, Se; R₁-R₃ = H, halogen, (un)substituted alkyl, (un)substituted alkenyl, (un)substituted alkynyl, (un)substituted cycloalkyl, (un)substituted Ph, (un)substituted heteroaryl; k = 0-6; l = 0-2; x = 0, 1; yr = 1-4] [e.g., (Bu)₄N⁺-N(CF₃)₂], useful as chem. intermediates and as liq. crystal

precursors (no data), are prep'd. by the reaction of alkali metal amine salts D+-N(CF₃)₂ (D = Group IA metal) [e.g., RbN(CF₃)₂] with salts [[R₁(CR₂R₃)_k]_lA_x]yK⁺-E (E = F, Cl, Br, I, BF₄, ClO₄, AsF₆, SbF₆, PF₆) (e.g., tetrabutylammonium tetrafluoroborate) in polar org. solvents (e.g., acetonitrile).

L9 ANSWER 7 OF 8 CAPLUS COPYRIGHT 2008 ACS on STN

Full Text

AN 2000:504821 CAPLUS
 DN 133:107335
 TI Carbon electrodes for double-layer capacitors I. Relations between ion and pore dimensions
 AU Salitra, Gregory; Soffer, Abraham; Eliad, Linoam; Cohen, Yair; Aurbach, Doron
 CS Department of Chemistry, Bar-Ilan University, Ramat-Gan, 52900, Israel
 SO Journal of the Electrochemical Society (2000), 147(7), 2486-2493
 CODEN: JESOAN; ISSN: 0013-4651
 PB Electrochemical Society
 DT Journal
 LA English
 AB We characterized activated carbon electrodes for elec. double-layer capacitor (EDLC) systems. High-surface-area carbons were prep'd. by carbonization of cotton cloth at elevated temps. (up to 1050°C), followed by activation at 900°C by oxidn. with CO₂ during different time periods. Sp. surface areas and characteristic pore sizes obtained from gas adsorption isotherms were correlated with those obtained from ion electroadsorption at the elec. double layer. Electrolytes studied included aq. LiCl, NaCl, and KCl solns. and nonaq. propylene carbonate solns. with LiBF₄ and (C₂H₅)₄NBF₄ salts. We found clear evidence that the porous carbons thus formed exhibit ion sieving properties, and that increasing activation time systematically increases the av. pore sizes of these carbons. The elec. double layer (EDL) capacity of these samples (calcd. from voltammetric measurements) depends strongly on the adsorption interaction of the ions in the pores, and hence the relationship between the av. pore size and the effective ion size dets. the specific EDL capacitance of these samples. The following order of dimension of adsorbed species was found, based on the ion sieving of the various synthesized carbons of different av. pore size--N₂, Na⁺(aq), Cl⁻(≈3.6 Å) < BF₄⁻ < TEA⁺(PC) < Li⁺(PC).

RE.CNT 27 THERE ARE 27 CITED REFERENCES AVAILABLE FOR THIS RECORD
 ALL CITATIONS AVAILABLE IN THE RE FORMAT

L9 ANSWER 8 OF 8 CAPLUS COPYRIGHT 2008 ACS on STN

Full Text

AN 1987:42927 CAPLUS
 DN 106:42927
 OREF 106:6993a,6996a
 TI Fluoroperoxide compounds of boron
 AU Ippolitov, E. G.; Chernyshov, B. N.; Shchetinina, G. P.; Brovkina, O. V.; Martynyuk, Yu. L.; Gorin, Yu. V.
 CS Inst. Vulkanol., Petropavlovsk-Kamchatskii, USSR
 SO Ukrainskii Khimicheskii Zhurnal (Russian Edition) (1986), 52(8), 818-23
 CODEN: UKZHAU; ISSN: 0041-6045
 DT Journal
 LA Russian
 AB 11B and 19F NMR study of M₂B₄O₇·xH₂O-MF-H₂O₂ solns. (M = NH₄⁺, Li, Na, K) indicated the formation of BF₄⁻, [BF(OOH)₃]⁻, [BF₂(OOH)₂]⁻, [BF₃(OOH)]⁻, [BF₃OH]⁻, [B₂F₄(O₂)₂]²⁻, [B₂F₂O₈H₂]²⁻, and [B(OOH)₄]⁻. An increase in H₂O₂ concn. increased the amt. of [B(OOH)₄]⁻ and [BF(OOH)₃]⁻. The change in compn. of the products was studied with increase of F⁻ concn. and an increase in time.

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FILE 'CAPLUS' ENTERED AT 07:50:40 ON 04 AUG 2008

L1 2 S TETRAFLUORBORATE (S) (METAL OR LI OR LITHIUM OR NA OR SODIUM
L2 5225 S TETRAFLUOROBORATE (S) (METAL OR LI OR LITHIUM OR NA OR SODIUM
L3 2137 S BF4 (S) (METAL OR LI OR LITHIUM OR NA OR SODIUM OR K OR POTAS
L4 7 S ?AMMONIUM (A) (BF4 OR TETRAFLUOROBORATE)
L5 3706 S ?AMMONIUM (A) (BF4 OR TETRAFLUOROBORATE)
L6 334 S (L1 OR L2) AND (L3 OR L4)
L7 128703 S ?AMMONIUM (A) (HALIDE OR FLUORIDE OR BROMIDE OR CHLORIDE OR I
L8 413 S L5 AND L7
L9 8 S L6 AND L7

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